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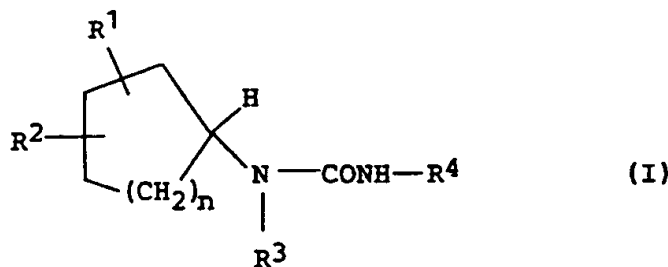
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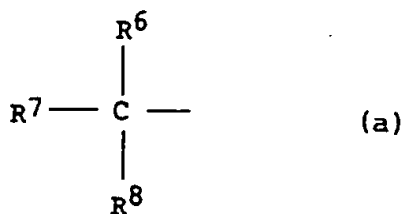
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 Cycloalkylurea compounds.

 A cycloalkylurea compound of the formula (I):



wherein R¹ is a group of the formula (a):

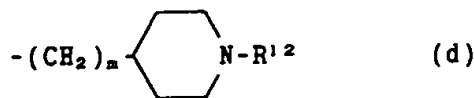
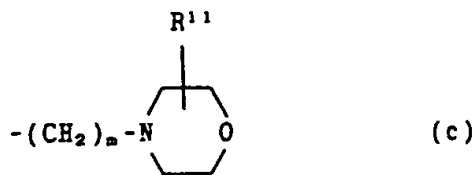


(wherein R^6 , R^7 and R^8 are the same or different and each is hydrogen or alkyl having 1 to 8 carbon atoms, with the proviso that compounds wherein two or three of R^6 , R^7 and R^8 are hydrogens are excluded), haloalkyl, cycloalkyl, alkoxy having 1 to 5 carbon atoms, phenyl, aralkyl or heteroaryl, or phenyl, aralkyl or heteroaryl each substituted on the aromatic ring by 1 to 3 substituents selected from alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 8 carbon atoms, cycloalkyl, cycloalkyloxy, haloalkyl, haloalkoxy, halogen, nitro, amino and substituted amino;

R^2 is hydrogen, phenyl or phenyl substituted by 1 to 3 substituents selected from alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, haloalkyl and halogen, wherein R^1 and R^2 may be substituted on the same carbon atom of cycloalkyl ring;

R^1 and R^2 together may form a substituted or unsubstituted cyclic hydrocarbon having 3 to 7 carbon atoms or a substituted or unsubstituted spiran;

R^3 is hydrogen, alkyl having 1 to 8 carbon atoms, cycloalkyl, cycloalkylalkyl, alkoxyalkyl, alkylthioalkyl, aralkyl substituted by alkylenedioxy, aralkyl, aralkyloxyalkyl, aralkyl or aralkyloxyalkyl each substituted on the aromatic ring by 1 to 3 substituents selected from alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 5 carbon atoms, halogen, nitro, hydroxy, amino, substituted amino, haloalkyl, alkylthio, benzyloxy and benzylthio, heteroarylalkyl, phenoxyalkyl, or heteroarylalkyl or phenoxyalkyl each substituted on the aromatic ring by 1 to 3 substituents selected from halogen, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, nitro, amino and haloalkyl, or a group of the formula (c) or (d):



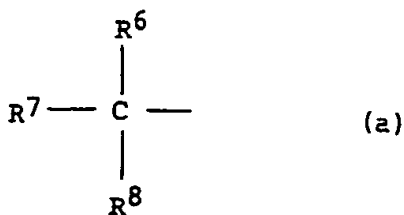
wherein m is 1 or 2, R^{11} is hydrogen or alkyl having 1 to 4 carbon atoms, R^{12} is hydrogen, alkyl having 1 to 4 carbon atoms or aralkyl;

R^4 is phenyl substituted by 1 to 3 substituents selected from alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, haloalkyl, halogen, amino and substituted amino;

n is 1, 2 or 3;

or a pharmaceutically acceptable salt thereof,

which compound inhibits ACAT and is useful as hypolipidemic and anti-atherosclerotic medicine.



10 (wherein R^6 , R^7 and R^8 are the same or different and each is hydrogen or alkyl having 1 to 8 carbon atoms, with the proviso that compounds wherein two or three of R^6 , R^7 and R^8 are hydrogens are excluded), haloalkyl, cycloalkyl, alkoxy having 1 to 5 carbon atoms, phenyl, aralkyl or heteroaryl, or phenyl, aralkyl or heteroaryl each substituted on the aromatic ring by 1 to 3 substituents selected from

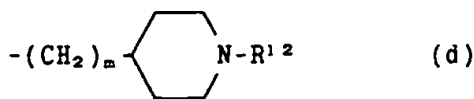
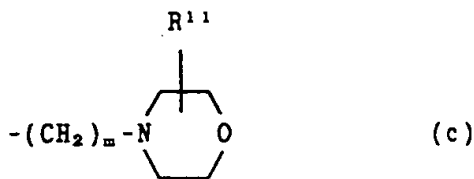
15 alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 8 carbon atoms, cycloalkyl, cycloalkyloxy, haloalkyl, haloalkoxy, halogen, nitro, amino and substituted amino;

R^2 is hydrogen, phenyl or phenyl substituted by 1 to 3 substituents selected from alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, haloalkyl and halogen, whererin R^1 and R^2 may be substituted on the same carbon atom of cycloalkyl ring;

20 R^1 and R^2 together may form a substituted or unsubstituted cyclic hydrocarbon having 3 to 7 carbon atoms or a substituted or unsubstituted spiran;

R^3 is hydrogen, alkyl having 1 to 8 carbon atoms, cycloalkyl, cycloalkylalkyl, alkoxyalkyl, alkylthioalkyl, aralkyl substituted by alkylendioxy, aralkyl, aralkyloxyalkyl, aralkyl or aralkyloxyalkyl each substituted on the aromatic ring by 1 to 3 substituents selected from alkyl having 1 to 4 carbon atoms, alkoxy

25 having 1 to 5 carbon atoms, halogen, nitro, hydroxy, amino, substituted amino, haloalkyl, alkylthio, benzyloxy and benzylthio, heteroarylalkyl, phenoxyalkyl, or heteroarylalkyl or phenoxyalkyl each substituted on the aromatic ring by 1 to 3 substituents selected from halogen, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, nitro, amino and haloalkyl, or a group of the formula (c) or (d):



45 wherein m is 1 or 2, R^{11} is hydrogen or alkyl having 1 to 4 carbon atoms, R^{12} is hydrogen, alkyl having 1 to 4 carbon atoms or aralkyl;

R^4 is phenyl substituted by 1 to 3 substituents selected from alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, haloalkyl, halogen, amino and substituted amino;

n is 1, 2 or 3;

50 or a pharmaceutically acceptable salt thereof.

2. A compound as disclosed in the above-mentioned 1 wherein R^1 is phenyl or phenyl substituted by 1 to 3 substituents selected from alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 8 carbon atoms, haloalkyl, halogen, nitro, amino and substituted amino; R^2 is phenyl or phenyl substituted by 1 to 3 substituents selected from alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, trifluoromethyl and halogen, whererin R^1 and R^2 may be substituted on the same carbon atom of cycloalkyl ring; or R^1 and R^2 together may form a substituted or unsubstituted cyclic hydrocarbon having 3 to 7 carbon atoms or a substituted or unsubstituted spiran; R^3 is hydrogen, alkyl having 1 to 8 carbon atoms, cycloalkyl, cycloalkylalkyl, aralkyl substituted by alkylendioxy, aralkyl, aralkyl substituted on the

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aromatic ring by 1 to 3 substituents selected from alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 5 carbon atoms, halogen, nitro, hydroxy, haloalkyl, alkylthio, benzyloxy and benzylthio, or heteroarylalkyl; R⁴ is phenyl substituted by 1 to 3 substituents selected from alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, trifluoromethyl, halogen and substituted amino; n is 2;

5 or a pharmaceutically acceptable salt thereof.

3. A compound as disclosed in the above-mentioned 1 wherein R¹ is secondary or tertiary alkyl having 3 to 5 carbon atoms, cycloalkyl or phenyl; R² is hydrogen; R³ is hydrogen, alkyl having 1 to 8 carbon atoms, cycloalkyl, cycloalkylalkyl, aralkyl substituted by alkylendioxy, aralkyl, aralkyl substituted on the aromatic ring by 1 to 3 substituents selected from alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 5 carbon atoms, halogen, nitro, hydroxy, haloalkyl, alkylthio, benzyloxy and benzylthio, or heteroarylalkyl; R⁴ is phenyl substituted by 1 to 3 substituents selected from alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms and halogen; n is 2; or a pharmaceutically acceptable salt thereof.

4. In detail, a compound as disclosed in the above-mentioned 1 selected from the group consisting of

15 N-(2,6-diisopropylphenyl)-N'-(2-phenylcyclohexyl)urea,
 N-(2,6-diisopropylphenyl)-N'-heptyl-N'-(4-phenylcyclohexyl)urea,
 trans-N-(2,6-diisopropylphenyl)-N'-benzyl-N'-(4-phenylcyclohexyl)urea,
 N-(2,6-diethylphenyl)-N'-benzyl-N'-(4-phenylcyclohexyl)urea,
 N-(2,4-difluorophenyl)-N'-benzyl-N'-(4-phenylcyclohexyl)urea,
 N-(2,6-diisopropylphenyl)-N'-(3,4-dimethoxybenzyl)-N'-(4-phenylcyclohexyl)urea,
 20 N-(2,4,6-trimethoxyphenyl)-N'-benzyl-N'-(4-phenylcyclohexyl)urea,
 N-(2,6-diisopropylphenyl)-N'-benzyl-N'-(4-tert-butylcyclohexyl)urea,
 trans-N-(2,6-diisopropylphenyl)-N'-(2-methylbenzyl)-N'-(4-phenylcyclohexyl)urea,
 trans-N-(2,6-diisopropylphenyl)-N'-(2-chlorobenzyl)-N'-(4-phenylcyclohexyl)urea,
 N-(2,6-diisopropylphenyl)-N'-(2-phenylethyl)-N'-(4-phenylcyclohexyl)urea,
 25 trans-N-(2,6-diisopropylphenyl)-N'-(3-phenylpropyl)-N'-(4-phenylcyclohexyl)urea,
 trans-N-(2,6-diisopropylphenyl)-N'-(1,1'-bicyclohexyl-4-yl)urea,
 trans-N-(2,6-diethylphenyl)-N'-cyclohexylmethyl-N'-(4-phenylcyclohexyl)urea,
 trans-N-(2,6-diisopropylphenyl)-N'-cyclohexylmethyl-N'-(4-phenylcyclohexyl)urea,
 N-(2,6-diisopropylphenyl)-N'-thenyl-N'-(4-phenylcyclohexyl)urea,
 30 trans-N-(2,6-diisopropylphenyl)-N'-furfuryl-N'-(4-phenylcyclohexyl)urea,
 N-(2,6-diethylphenyl)-N'-(2-methylbenzyl)-N'-(4-phenylcyclohexyl)urea,
 N-(2,6-diethylphenyl)-N'-(2-chlorobenzyl)-N'-(4-phenylcyclohexyl)urea and
 trans-N-(2,6-diisopropylphenyl)-N'-(2-pyridylmethyl)-N'-(4-phenylcyclohexyl)urea

or a pharmaceutically acceptable salt thereof.

5. Preferably, a compound as disclosed in the above-mentioned 1 selected from the group consisting of

35 trans-N-(2,6-diisopropylphenyl)-N'-benzyl-N'-(4-phenylcyclohexyl)urea,
 trans-N-(2,6-diisopropylphenyl)-N'-(2-methylbenzyl)-N'-(4-phenylcyclohexyl)urea,
 trans-N-(2,6-diisopropylphenyl)-N'-(2-chlorobenzyl)-N'-(4-phenylcyclohexyl)urea and
 trans-N-(2,6-diethylphenyl)-N'-cyclohexylmethyl-N'-(4-phenylcyclohexyl)urea,

40 or a pharmaceutically acceptable salt thereof.

6. Also preferably, a compound as disclosed in the above-mentioned 1 which is N-(2,6-diisopropylphenyl)-N'-(4,4-diphenylcyclohexyl)urea, or a pharmaceutically acceptable salt thereof.

7. A pharmaceutical composition consisting of an effective amount of the compound as disclosed in the above-mentioned 1 or a pharmaceutically acceptable salt thereof and pharmaceutical carriers.

45 The definitions of each symbols in the formula (I) are exemplified as follows:

In R¹, alkyl having 1 to 8 carbon atoms means, for example, methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, tert-butyl, pentyl, hexyl, heptyl, octyl, 1-ethylpropyl or 2,2-dimethylpropyl; the group of formula (a) in R¹ means preferably secondary or tertiary alkyl having 3 to 12 carbon atoms and is exemplified by, for example, isopropyl, 1-methylpropyl, 1-methylbutyl, 1-ethylpropyl, tert-butyl, tert-pentyl,
 50 1-methylhexyl, 1-methylheptyl, 1-methyloctyl, 1-methylnonyl, 1-ethylnonyl, 1,1-dimethylhexyl or 1,1-dimethylnonyl; haloalkyl has 1 to 8 carbon atoms in the alkyl moiety and is exemplified by, for example, fluoromethyl, trifluoromethyl, chloroethyl, 2,2,2-trifluoroethyl, perfluoroethyl, chloropropyl, chlorobutyl, chloropentyl, chlorohexyl or chloroheptyl; cycloalkyl has 3 to 7 carbon atoms and is exemplified by, for example, cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl or cycloheptyl, preferably cyclohexyl; alkyl having
 55 1 to 4 carbon atoms means, for example, methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl or tert-butyl; alkoxy having 1 to 4 carbon atoms means, for example, methoxy, ethoxy, propoxy, isopropoxy, butoxy, isobutoxy, sec-butoxy or tert-butoxy; alkoxy having 5 carbon atoms means pentyloxy, isopentyloxy or tert-pentyloxy; alkoxy having 1 to 8 carbon atoms means hexyloxy, heptyloxy or octyloxy in addition to

the alkoxy having 1 to 5 carbon atoms; haloalkoxy has 1 to 4 carbon atoms and is exemplified by, for example, fluoromethoxy, trifluoromethoxy, chloroethoxy, 2,2,2-trifluoroethoxy, perfluoroethoxy, chloropropoxy or chlorobutoxy; cycloalkyloxy has 3 to 7 carbon atoms in the cycloalkyl moiety and is exemplified by, for example, cyclopropyloxy, cyclobutyloxy, cyclopentyloxy, cyclohexyloxy or cycloheptyloxy; halogen means fluorine, chlorine, bromine or iodine; substituted amino means mono- or dialkylamino (e.g. methylamino, dimethylamino, ethylamino, diethylamino, propylamino, dipropylamino), acylamino (e.g. formylamino, acetylamino, propionylamino, butyrylamino, isobutyrylamino, benzoylamino), alkoxycarbonylamino (e.g. methoxycarbonylamino, ethoxycarbonylamino, propoxycarbonylamino), aralkyloxycarbonylamino (e.g. benzyloxycarbonylamino, 2-phenylethyloxycarbonylamino) or cyclic amino (e.g. pyrrolidinyl, piperidino, morpholino, thiomorpholino, 1-piperazinyl, 4-methyl-1-piperazinyl); aralkyl means, for example, benzyl, 1-phenylethyl, 2-phenylethyl, 1-phenylpropyl, 2-phenylpropyl, 3-phenylpropyl, 2-methyl-2-phenylpropyl, 4-phenylbutyl, 5-phenylpentyl, 1-naphthylmethyl or 1-(1-naphthyl)ethyl; heteroaryl means, for example, thienyl, pyridyl, furyl, pyrrolyl, oxazolyl, thiazolyl, imidazolyl, pyrazolyl, pyrimidinyl or indolyl.

Preferred groups of R¹ are secondary or tertiary alkyl having 3 to 5 carbon atoms (e.g. isopropyl, 1-methylpropyl, 1-methylbutyl, 1-ethylpropyl, tert-butyl, tert-pentyl), cycloalkyl (e.g. cyclohexyl) and phenyl.

In R², alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, haloalkyl and halogen which are substituents of phenyl are exemplified by those mentioned in R¹. Preferred groups of R² are hydrogen, phenyl and phenyl substituted by halogen.

Substituted or unsubstituted cyclic hydrocarbon having 3 to 7 carbon atoms formed by R¹ and R² means, for example, cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, methylcyclopentyl, methylcyclohexyl, ethylcyclohexyl or butylcyclohexyl, preferably cyclohexyl; substituted or unsubstituted spiran formed by R¹ and R² with cycloalkyl ring means, for example, spiro[5.5]undec-3-yl, spiro[4.5]dec-8-yl, spiro[4.5]dec-2-yl, spiro[2.5]oct-6-yl or spiro[4.5]dec-6-yl; the substituent(s) for cyclic hydrocarbon and spiran is(are) selected from alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, halogen, nitro and amino.

In R³, alkyl having 1 to 8 carbon atoms, cycloalkyl and aralkyl are exemplified by those mentioned in R¹; cycloalkylalkyl has 3 to 7 carbon atoms in the cycloalkyl moiety and 1 to 4 carbon atoms in the alkyl moiety and is exemplified by, for example, cyclopropylmethyl, cyclopentylmethyl, cyclohexylmethyl, cycloheptylmethyl, 2-cyclohexylethyl or 3-cyclohexylpropyl; alkoxyalkyl has 1 to 4 carbon atoms in the alkoxy moiety and 1 to 4 carbon atoms in the alkyl moiety and is exemplified by, for example, methoxymethyl, methoxyethyl, methoxypropyl, methoxybutyl, ethoxymethyl, ethoxyethyl or ethoxypropyl; alkylthioalkyl has 1 to 4 carbon atoms in the alkylthio moiety and 1 to 4 carbon atoms in the alkyl moiety and is exemplified by, for example, methylthiomethyl, methylthioethyl, methylthiopropyl, methylthiobutyl, ethylthiomethyl, ethylthioethyl or ethylthiopropyl; aralkyl substituted by alkylendioxy means (1,3-benzodioxol-5-yl)methyl or (1,4-benzodioxan-6-yl)methyl; aralkyloxyalkyl has 1 to 4 carbon atoms in the alkyl moiety and is exemplified by, for example, benzyloxymethyl, 2-benzyloxyethyl, 3-benzyloxypropyl, 2-phenylethoxymethyl or 2-(2-phenylethoxy)ethyl; heteroarylalkyl has 1 to 4 carbon atoms in the alkyl moiety and is exemplified by, for example, thenyl, 2-, 3- or 4-pyridylmethyl, 2-(2-pyridyl)ethyl, furfuryl, 2-(2-furyl)ethyl or 2-pyrimidinylmethyl; phenoxyalkyl has 1 to 4 carbon atoms in the alkyl moiety and is exemplified by, for example, phenoxyethyl, 2-phenoxyethyl, 3-phenoxypropyl or 4-phenoxybutyl.

With regard to substituents in R³, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 5 carbon atoms, halogen, substituted amino and haloalkyl are exemplified by those mentioned in R¹; alkylthio has 1 to 4 carbon atoms in the alkyl moiety and is exemplified by, for example, methylthio, ethylthio, propylthio, isopropylthio or butylthio; alkyl having 1 to 4 carbon atoms and aralkyl which are represented by R¹¹ or R¹² in the formulas (c) and (d) are exemplified by those mentioned in R¹.

Preferred groups of R³ are hydrogen, alkyl having 1 to 8 carbon atoms, cycloalkyl, cycloalkylalkyl, aralkyl substituted by alkylendioxy, aralkyl, aralkyl substituted on the aromatic ring by 1 to 3 substituents selected from alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 5 carbon atoms, halogen, nitro, hydroxy, haloalkyl, alkylthio, benzyloxy and benzylthio, or heteroarylalkyl.

More preferred groups of R³ are cycloalkyl (e.g. cyclohexyl), cycloalkylalkyl (e.g. cyclohexylmethyl), aralkyl (e.g. benzyl, 2-phenylethyl, 3-phenylpropyl, 4-phenylbutyl) and aralkyl substituted on the aromatic ring by substituent(s) (e.g. 2-chlorobenzyl, 2-fluorobenzyl, 2-methylbenzyl, 2-bromobenzyl, 2-methoxybenzyl, 2-ethoxybenzyl, 2-methylthiobenzyl, 2-(2-chlorophenyl)ethyl).

In R⁴, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, haloalkyl, halogen and substituted amino which are substituents of phenyl are exemplified by those mentioned in R¹.

Preferred groups of R⁴ are phenyl substituted by 1 to 3 substituents selected from alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms and halogen;

More preferred groups of R⁴ are 2,6-diethylphenyl, 2,6-diisopropylphenyl, 2,4-difluorophenyl and 2,4,6-

trimethoxyphenyl.

The pharmaceutically acceptable salts of the compounds of formula (I) include salts with an inorganic acid such as hydrochloric acid, hydrobromic acid, sulfuric acid etc., or with an organic acid such as acetic acid, fumaric acid, maleic acid, benzoic acid, citric acid, malic acid, methanesulfonic acid, benzenesulfonic acid etc. The present invention also includes hydrates and solvates of the compounds of formula (I).

In the compounds of formula (I), there exist compounds having cis- or trans-configuration between the urea group and the substituents: R¹ or R² regarding the cycloalkyl ring. The present invention embraces cis- and trans-isomers and the mixture thereof. When the compounds of the present invention have one or more asymmetric center, there exist a variety of optical isomers. The individual optical isomers, racemate thereof and diastereomer or the mixture thereof are encompassed in the present invention.

The compounds of the present invention can be prepared by, for example, the following methods outlined in Reaction Scheme.

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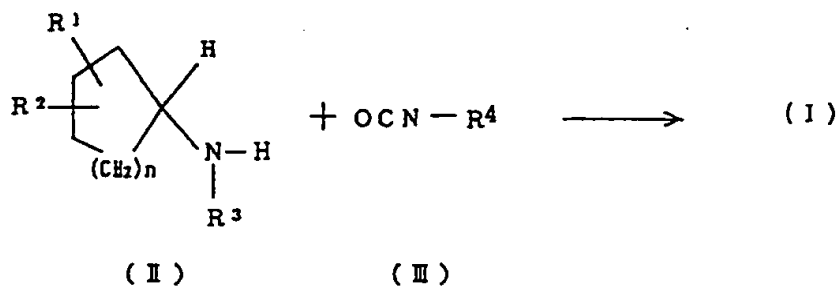
Reaction Scheme

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Method A

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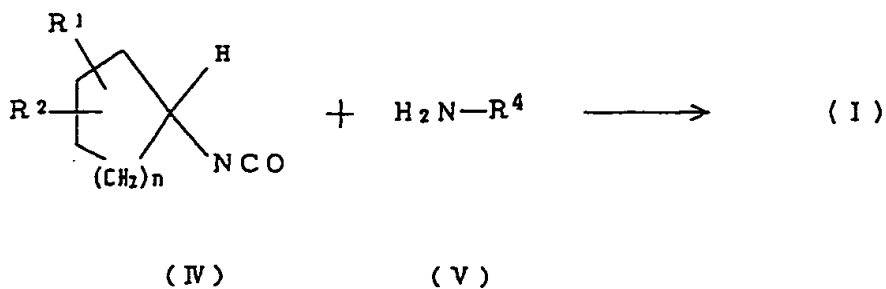


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Method B

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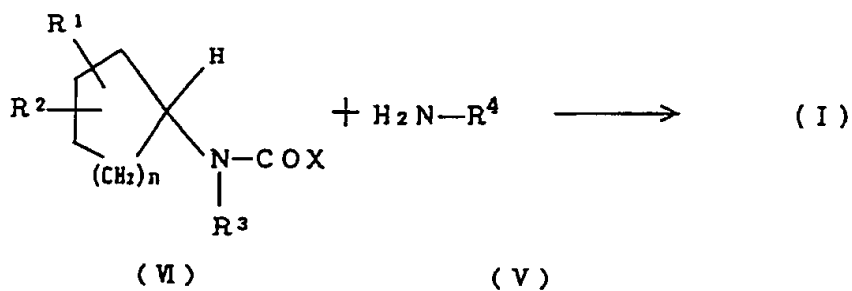


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Method C

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50 (In the above formulas, X is halogen, and other symbols are as defined above.)

55 Method A

The compounds of formula (I) can be synthesized by reacting a cycloalkylamine of the formula (II) or a salt thereof with an isocyanic ester of the formula (III) in an appropriate solvent. Any solvent inert to the

reaction can be used as the solvent. Thus, for example, use may be made of ethers such as diethyl ether, diisopropyl ether, dimethoxyethane or tetrahydrofuran; aromatic hydrocarbons such as benzene, toluene or xylene; esters such as methyl acetate or ethyl acetate; ketones such as acetone or methyl ethyl ketone; halogenated hydrocarbons such as dichloromethane, chloroform or dichloroethane; N,N-dimethylformamide; acetamide; acetonitrile; dimethyl sulfoxide and pyridine. The reaction may be carried out at any temperature between -20° C and 150° C, preferably between 0° C and 100° C. The reaction will be significantly proceeded in the presence of a base such as triethylamine, N-methylmorpholine, diisopropylethylamine, 1,4-diazabicyclo[2.2.2]octane, pyridine, picoline, 4-dimethylaminopyridine, 4-(1-pyrrolidinyl)pyridine or N,N-dimethylaniline.

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Method B

The compounds of formula (I) wherein R³ is hydrogen can be synthesized by reacting an isocyanic ester of the formula (IV) with an amine of the formula (V) under similar conditions as in Method A.

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Method C

The compounds of formula (I) wherein R³ is the substituent other than hydrogen can be synthesized by reacting a carbamoyl halide of the formula (VI) with an amine of the formula (V) under similar conditions as in Method A.

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The starting compounds, i.e. cycloalkylamine of the formula (II), can be synthesized by per se known methods.

The compounds of formula (I) can be converted into the abovementioned pharmaceutically acceptable salts by treating them with an inorganic or an organic acid in a conventional manner.

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Among the compounds of formula (I), there exist cis- and trans-isomers. They can be synthesized respectively by using cis- and trans-compounds of the formula (II), (IV) or (VI) as the starting compound. When the obtained compound is a mixture thereof, it can be separated into cis- and trans-isomers by means of a conventional method such as recrystallization or chromatography. The compounds of the present invention having a asymmetric center are usually obtained as racemates. The racemates can be resolved into the optical isomers by a conventional method such as fractional recrystallization or chromatography, or each optical isomer can be produced by employing the optically active starting compounds. The compounds having two or more asymmetric centers are usually obtained as each diastereomers or a mixture thereof. Each diastereomers can be resolved by means of fractional recrystallization or chromatography.

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The following pharmacological experiments will illustrate that the compounds of the present invention have potent ACAT inhibitory activity and hypolipidemic activity.

Pharmacological Experiment 1: ACAT inhibitory activity

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ACAT activity was determined according to the method of Heider et al. [J. Lipid Res., 24, 1127 (1983)]. Microsomes from intestinal mucosa of male Japanese White rabbits fed with 2% cholesterol-supplemented chow were used as the enzyme. The mixture of 0.154 M potassium phosphate buffer (pH 7.4), containing [¹⁴C]oleoyl-CoA, bovine serum albumin and test compounds were preincubated at 37° C for 5 min before the addition of the enzyme. The mixture was incubated for 3 min. The reaction was stopped by the addition of the mixture of chloroform and methanol (2:1). After centrifugation, the chloroform phase was collected and dried. The residue was dissolved in n-hexane and spotted onto silica gel plates for thin layer chromatography (TLC). After the plates were developed, radioactivity in cholesteryl ester fraction was measured by using radio-TLC analyzer.

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As a result, the compounds of the present invention have been found to have potent ACAT inhibitory activity with IC₅₀ values ranging from about 10 nM to about 50 nM.

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Pharmacological Experiment 2: Cholesterol-lowering activity

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Male Lewis rats were fed a diet supplemented with 1.5% cholesterol, 0.5% cholic acid and 10% coconut oil. Test compounds were added into a diet. After 3 days, total cholesterol in serum and liver was determined by enzymatic assay.

As a result, the test compounds caused significant decrease in serum and liver cholesterol levels in the cholesterol-loaded rats compared with those of control group. The same tests were conducted using mice

and rabbits and the results similar to those in rats were observed.

As is clear from the foregoing pharmacological experiments, the compounds of the present invention exhibit potent ACAT inhibitory activity and cause significant decrease in serum and liver cholesterol levels in hyperlipidemic animals such as rats, mice or rabbits.

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Pharmacological Experiment 3: Inhibition of cholesterol esterification in cultured macrophages

Cholesterol esterification activity in macrophages, which has been known to be stimulated by modified LDL, was determined according to the method of Brown et al. (J. Biol. Chem., 255, 9344 (1980)). The
 10 peritoneal macrophages from male ddY mice were preincubated with test compounds at 37° C for 2 hours under 5% CO₂/95% air. They were then incubated with [1-¹⁴C]oleic acid, bovine serum albumine and acetylated LDL for 5 hours. The reaction was stopped and the intracellular lipids were extracted with hexane-propanol (3:2). Radioactivity in cholesteryl ester was measured as described in the Experiment 1. As
 15 a result, the compounds of the present invention have been found to be potent inhibitors of cholesterol esterification in macrophages. IC₅₀ values of the compounds were from 10⁻⁷ to 10⁻⁹ M.

Pharmacological Experiment 4: Acute toxicity

The compounds of Examples 53 and 78 were orally administered to six male mice at the dosage of
 20 1000 mg/kg, and no death occurrences were observed for 7 days.

The compounds of the present invention are potent ACAT inhibitors. They suppress the formation of cholesteryl esters in cultured macrophages and some other cells. They decrease serum and liver cholesterol levels in hyperlipidemic animals and are expected to suppress the absorption of cholesterol at intestinal mucosa and decrease the serum cholesterol level in human. They are also expected to suppress
 25 the cholesteryl ester formation in macrophages and prevent the accumulation of lipids in the artery wall. It has been shown that they have advantageous characters, concerning absorption through oral administration, bioavailability, duration time and safety. Therefore new hypolipidemic and anti-atherosclerotic medicines with high safety and potent activity would be obtained through the present invention.

The compounds of the present invention can be safely administered orally or parenterally to human
 30 beings in the form of pharmaceutical compositions such as powders, tablets, capsules, granules, suppositories or injectable solutions. The pharmaceutical composition can be prepared by mixing a therapeutically effective amount of the compound with a pharmaceutically acceptable additives such as carrier, excipient or diluent. The dose may vary depending upon the body weight, the severity of the patient to be treated or the age of the patient, but the daily dose for human adults preferably ranges 10 mg to 500
 35 mg in single or multiple dose.

EXAMPLE

The present invention will be explained in more detail by the following examples, but these are not to
 40 be construed as limiting the present invention.

Example 1

To a solution of 1.6 g of 4-tert-butylcyclohexylamine in 15 ml of dichloromethane was added a solution
 45 of 2.2 g of 2,6-diisopropylphenylisocyanate in dichloromethane dropwise in the presence of pyridine at 0° C. Thereafter the mixture was stirred for 1.5 hours at room temperature and concentrated. The residue was partitioned between isopropyl ether and water, and the organic layer was successively washed with 0.5N hydrochloric acid, water and sodium sulfate solution. The organic layer was dried over magnesium sulfate and then distilled away. The resulting residue was crystallized from ethyl acetate to give 2.2 g of N-(2,6-
 50 diisopropylphenyl)-N'-(4-tert-butylcyclohexyl)urea, melting at 270° C.

Example 2

The reaction and procedure were conducted in the same manner as in Example 1 using 1.0 g of trans-
 55 4-tert-butylcyclohexylamine to give 0.5 g of trans-N-(2,6-diisopropylphenyl)-N'-(4-tert-butylcyclohexyl)urea, melting at 264-266° C.

Example 3

The reaction and procedure were conducted in the same manner as in Example 1 using cis-4-tert-butylcyclohexylamine to give cis-N-(2,6-diisopropylphenyl)-N'-(4-tert-butylcyclohexyl)urea, melting at 179-180.5° C.

5 Example 4

To a solution of 1 g of 4-phenylcyclohexylamine in 10 ml of dichloromethane was added 2,6-diethylphenylisocyanate dropwise in the presence of pyridine and the mixture was stirred for 2 hours at room temperature. To the resultant mixture was added 30 ml of diluted hydrochloric acid and the solution
10 was extracted with chloroform. The extract was washed with water, aqueous sodium hydrogencarbonate solution and saturated brine, and then dried over magnesium sulfate. The solvent was distilled away and the residue was crystallized from methanol to give 0.4 g of N-(2,6-diethylphenyl)-N'-(4-phenylcyclohexyl)urea, melting at 247-249° C.

15 Example 5

The reaction and procedure was conducted in the same manner as in Example 1 using 1.5 g of 2-phenylcyclohexylamine to give 1.2 g of N-(2,6-diisopropylphenyl)-N'-(2-phenylcyclohexyl)urea, melting at 225-226° C.

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Example 6

To a solution of 1.6 g of 4-isopropylcyclohexylisocyanate in chloroform is added a solution of 1.8 g of 2,4,6-trimethoxyaniline in chloroform dropwise in the presence of pyridine. The mixture is stirred at room
25 temperature until starting compounds disappear on TLC. The resultant mixture is concentrated and the residue is dissolved in ethyl acetate. The solution is washed with diluted hydrochloric acid, aqueous sodium hydrogencarbonate solution and brine. The organic layer is dried over magnesium sulfate and filtered off. The filtrate is distilled away and the resulting residue is crystallized to give N-(2,4,6-trimethoxyphenyl)-N'-(4-isopropylcyclohexyl)urea.

30

Example 7

To a solution of 0.82 g of 2-tert-butyl-6-methylaniline in pyridine is added a solution of 1.2 g of N-methyl-N-(2-phenylcyclohexyl)-carbamoylchloride in dichloromethane dropwise. The mixture is stirred at
35 room temperature until starting compounds disappear on TLC. The mixture is concentrated and the residue is dissolved in ethyl ether. The solution is washed with diluted hydrochloric acid, aqueous sodium hydrogencarbonate solution and brine. The organic layer is dried over sodium sulfate and filtered off. The filtrate is distilled away and the resulting residue is crystallized to give N-(2-tert-butyl-6-methylphenyl)-N'-methyl-N-(2-phenylcyclohexyl)urea.

40

The following compounds can be prepared in a similar manner as the above examples.

- (8) N-(2,4-difluorophenyl)-N'-(4,4-bis(4-chlorophenyl)cyclohexyl)-urea, melting at 190-191° C
 (9) N-(2,4-dimethoxyphenyl)-N'-(4,4-bis(4-chlorophenyl)cyclohexyl)-urea, melting at 145-147° C
 (10) N-(2,4-difluorophenyl)-N'-(4,4-diphenylcyclohexyl)urea, melting at 250-252° C
 (11) N-(3-methylphenyl)-N'-(4,4-bis(4-chlorophenyl)cyclohexyl)-urea, melting at 150-152° C
 45 (12) N-(3-trifluoromethylphenyl)-N'-(4,4-bis(4-chlorophenyl)cyclohexyl)urea, melting at 178-180° C
 (13) N-(3-methoxyphenyl)-N'-(4,4-bis(4-chlorophenyl)cyclohexyl)-urea, melting at 138-140° C
 (14) N-(4-bromophenyl)-N'-(4,4-bis(4-chlorophenyl)cyclohexyl)-urea, melting at 199-202° C
 (15) N-(2-ethylphenyl)-N'-(4,4-bis(4-chlorophenyl)cyclohexyl)-urea, melting at 150-152° C
 (16) N-(2,4-dichlorophenyl)-N'-(4,4-diphenylcyclohexyl)urea, melting at 235-237° C
 50 (17) N-(2,4-difluorophenyl)-N'-heptyl-N'-(4,4-bis(4-chlorophenyl)cyclohexyl)urea, melting at 86-90° C
 (18) N-(3-trifluoromethylphenyl)-N'-(4,4-bis(4-chlorophenyl)cyclohexyl)-N'-heptylurea, melting at 138-140° C
 (19) N-(3-trifluoromethylphenyl)-N'-(4,4-diphenylcyclohexyl)urea, melting at 236-238° C
 (20) N-(2,6-diisopropylphenyl)-N'-(4,4-bis(4-chlorophenyl)cyclohexyl)urea, melting at 255-257° C
 55 (21) N-(2,6-diisopropylphenyl)-N'-(decahydro-2-naphthyl)urea, melting at 229° C
 (22) N-(2,6-diisopropylphenyl)-N'-(4-phenylcyclohexyl)urea, melting at 195-196° C
 (23) N-(2,6-diethylphenyl)-N'-(4,4-diphenylcyclohexyl)urea, melting at 206-207° C
 (24) N-(2,6-diisopropylphenyl)-N'-(4,4-diphenylcyclohexyl)urea, melting at 243-244° C

- (25) N-(2,6-diisopropylphenyl)-N'-(3-phenylcyclohexyl)urea, melting at 214-215.5° C
 (26) N-(2,6-diethylphenyl)-N'-(2-phenylcyclohexyl)urea, melting at 249-250° C
 (27) N-(2,6-diisopropylphenyl)-N'-(4-tert-butylcyclohexyl)-N'-methylurea, melting at 256-258° C
 (28) N-(2,4-difluorophenyl)-N'-(4-tert-butylcyclohexyl)urea, melting at 225-227° C
 5 (29) N-(2,6-diisopropylphenyl)-N'-(spiro[5.5]undec-3-yl)urea, melting at 249.5° C
 (30) N-(2,6-diisopropylphenyl)-N'-(4-(4-pyridyl)cyclohexyl)urea 1/4 hydrate, melting at 276-277° C
 (31) N-(2,6-diisopropylphenyl)-N'-(4-tert-butylcyclohexyl)-N'-octylurea
 (32) N-(2,6-diisopropylphenyl)-N'-(spiro[2.5]oct-6-yl)urea
 (33) N-(2,6-diethylphenyl)-N'-(2-dicyclohexyl)urea
 10 (34) N-(2,6-diisopropylphenyl)-N'-(4-cyclopropylmethylcyclohexyl)urea
 (35) N-(2,6-diethylphenyl)-N'-(4-(2-phenylethyl)cyclohexyl)urea
 (36) N-(2-isopropyl-6-methylphenyl)-N'-(2-(2-thienyl)cyclopentyl)urea
 (37) N-(2,6-diisopropylphenyl)-N'-(2-tert-butylcyclohexyl)urea
 (38) N-(2,4,6-trimethoxyphenyl)-N'-(2-(4-isopropylphenyl)cyclohexyl)urea
 15 (39) N-(2,4,6-trimethylphenyl)-N'-(4-(1-ethylpropyl)cyclohexyl)urea
 (40) N-(4-dimethylaminophenyl)-N'-(3-(4-butoxyphenyl)cyclopentyl)urea
 (41) N-(4-acetyaminophenyl)-N'-(4-(4-nitrobenzyl)cycloheptyl)-N'-benzylurea
 (42) N-(2-tert-butyl-6-methylphenyl)-N'-(4-(1,1-dimethyloctyl)cyclohexyl)urea
 (43) N-(2,6-diisopropylphenyl)-N'-(4-tert-butylcyclohexyl)-N'-heptyl-urea, melting at 169-170° C
 20 (44) N-(2,6-diisopropylphenyl)-N'-(bicyclo[4.4.0]dec-2-yl)urea, melting at 217-218° C
 (45) N-(2,6-diisopropylphenyl)-N'-(4-(4-hexyloxyphenyl)cyclohexyl)-urea, melting at 225-226° C
 (46) N-(2,6-diisopropylphenyl)-N'-heptyl-N'-(4-phenylcyclohexyl)urea, melting at 143-144° C
 (47) N-(4-dimethylaminophenyl)-N'-(4-tert-butylcyclohexyl)urea, melting at 141-142° C
 (48) cis-N-(2,6-diisopropylphenyl)-N'-(2-phenylcyclohexyl)urea, melting at 187-189° C
 25 (49) N-(2,6-diisopropylphenyl)-N'-benzyl-N'-(4-phenylcyclohexyl)urea, melting at 144-146° C
 (50) N-(2,6-di-tert-butylphenyl)-N'-(4-tert-butylcyclohexyl)urea

Example 51

30 A solution of 5 g of 4-phenylcyclohexanone and cyclohexylmethylamine in 100 ml of benzene was refluxed for 2 hours with removing water and the solvent was distilled away in vacuo. The residue was dissolved in methanol and to the solution was added 1.5 g of sodium borohydride. The mixture was stirred at room temperature and the solvent was distilled away in vacuo. To the residue was added water and extracted with diethyl ether. The organic layer was washed with saturated brine and dried over magnesium sulfate. The solvent was distilled away to give 7 g of N-(4-phenylcyclohexyl)-N-cyclohexylmethylamine as
 35 an oil. (N-(4-phenylcyclohexyl)-N-cyclohexylmethylamine hydrochloride, melting at 267-270° C with decomposition)

To the solution of 2 g of N-(4-phenylcyclohexyl)-N-cyclohexylmethylamine in dichloromethane was added 2 ml of pyridine and to the mixture was added a solution of 1.6 g of 2,6-diethylphenylisocyanate in
 40 30 ml of dichloromethane dropwise. After the mixture was stirred at room temperature, to the solution was added diluted hydrochloric acid and extracted with ethyl acetate. The extract was washed with water, aqueous sodium hydrogencarbonate solution and saturated brine and then dried over magnesium sulfate. The solvent was distilled away in vacuo to give 2 g of crude crystals. The crystals were recrystallized from the mixture of ethyl acetate and hexane to give 1.2 g of N-(2,6-diethylphenyl)-N'-cyclohexylmethyl-N'-(4-
 45 phenylcyclohexyl)urea, melting at 166-168° C.

Example 52

A solution of 5 g of 4-phenylcyclohexanone and benzylamine in 100 ml of benzene was refluxed with
 50 removing water and the mixture was concentrated in vacuo. The residue was dissolved in tetrahydrofuran and to the solution was added the equivalent weight of sodium bis(2-methoxyethoxy)aluminum hydride (70% toluene solution) under ice cooling. After completion of the reaction, to the mixture was added ammonium chloride solution and extracted with diethyl ether. The organic layer was washed with saturated brine and dried over magnesium sulfate. The solvent was distilled away to give 6.5 g of oil. The oil was
 55 chromatographed on a silica gel column using a mixture of hexane and ethyl acetate (20:1) as an eluent to give 3.9 g of cis-N-benzyl-N-(4-phenylcyclohexyl)amine. (cis-N-benzyl-N-(4-phenylcyclohexyl)amine hydrochloride, melting at 208-210° C with decomposition)

To a solution of 400 mg of cis-N-benzyl-N-(4-phenylcyclohexyl)amine in 20 ml of dichloromethane were

added 2 ml of pyridine and then a solution of 310 mg of 2,6-diisopropylphenylisocyanate in 5 ml of dichloromethane dropwise. The mixture was stirred at room temperature, poured into diluted hydrochloric acid and extracted with ethyl acetate. The extract was washed with water, aqueous sodium hydrogencarbonate solution and saturated brine and then dried over magnesium sulfate. The solvent was distilled away
 5 in vacuo to give crude crystals. The crystals were recrystallised from the mixture of ethyl acetate and hexane to give 460 mg of cis-N-(2,6-diisopropylphenyl)-N'-benzyl-N'-(4-phenylcyclohexyl)urea, melting at 153-154° C.

Example 53

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A solution of 50 g of 4-phenylcyclohexanone and 30.7 g of benzylamine in 250 ml of benzene was refluxed for 3 hours with removing water and the solvent was distilled away in vacuo. The residue was dissolved in methanol and to the solution was added 10.8 g of sodium borohydride. The mixture was stirred at room temperature and the solvent was distilled away in vacuo. To the residue was added water and
 15 extracted with ethyl acetate. The organic layer was washed with water and saturated brine and dried over magnesium sulfate. The solvent was distilled away and ethyl acetate was added to the resulting oil. To the solution was added conc. hydrochloric acid and the precipitated crystals were collected by filtration. The crystals were recrystallized from the mixture of ethyl acetate and methanol to give 40.3 g of trans-N-benzyl-N-(4-phenylcyclohexyl)amine hydrochloride, melting at 244-248° C with decomposition.

20

75.4 g of trans-N-benzyl-N-(4-phenylcyclohexyl)amine hydrochloride was poured into aqueous sodium hydroxide solution and extracted with ethyl acetate. The extract was washed with water, dried and the solvent was distilled away. The residue was dissolved in 600 ml of dichloromethane and to the solution was added 60 ml of pyridine, and then added a solution of 51 g of 2,6-diisopropylphenylisocyanate in 100 ml of dichloromethane dropwise. After completion of the reaction, the solution was poured into diluted hydrochloric acid and extracted with ethyl acetate. The extract was washed with water, sodium hydrogencarbonate solution and saturated brine and then dried over magnesium sulfate. The solvent was distilled away in vacuo to give crude crystals. The crystals were recrystallized from the mixture of ethyl acetate and hexane to give
 25 73 g of trans-N-(2,6-diisopropylphenyl)-N'-benzyl-N'-(4-phenylcyclohexyl)urea, melting at 148-149° C.

30

- (54) N-(2,6-diisopropylphenyl)-N'-(1,1'-bicyclohexyl-4-yl)urea, melting at 259-261° C
- (55) N-(2,6-diethylphenyl)-N'-benzyl-N'-(4-phenylcyclohexyl)urea, melting at 148-150° C
- (56) N-(2,4-difluorophenyl)-N'-benzyl-N'-(4-phenylcyclohexyl)urea, melting at 140-142° C
- (57) N-(2,6-diisopropylphenyl)-N'-(1,1'-bicyclohexyl-2-yl)urea, melting at 226° C
- (58) N-(2,5-dimethoxyphenyl)-N'-(4-tert-butylcyclohexyl)urea, melting at 189-191° C
- (59) cis-N-(2,6-diisopropylphenyl)-N'-(2-phenylcyclohexyl)urea, melting at 138-140° C
- 35 (60) trans-N-(2,6-diisopropylphenyl)-N'-(3,4-dimethoxybenzyl)-N'-(4-phenylcyclohexyl)urea, melting at 129-130° C (as crystals containing 1 molecule of ethanol)
- (61) N-(2,4,6-trimethoxyphenyl)-N'-benzyl-N'-(4-phenylcyclohexyl)urea, melting at 188-190° C
- (62) N-(2,4,6-trimethoxyphenyl)-N'-(2-phenylcyclohexyl)urea, melting at 182-183° C
- (63) cis-N-(2,6-diisopropylphenyl)-N'-(1,1'-bicyclohexyl-4-yl)urea, melting at 208° C
- 40 (64) N-(2,6-diisopropylphenyl)-N'-benzyl-N'-(4-tert-butylcyclohexyl)urea, melting at 155-159° C
- (65) trans-N-(2,6-diisopropylphenyl)-N'-(2-methylbenzyl)-N'-(4-phenylcyclohexyl)urea, melting at 158-160° C
- (66) trans-N-(2,6-diisopropylphenyl)-N'-(2-chlorobenzyl)-N'-(4-phenylcyclohexyl)urea, melting at 197-198° C
- 45 (67) N-(2,6-diisopropylphenyl)-N'-(4-nitrobenzyl)-N'-(4-phenylcyclohexyl)urea, melting at 214-216° C
- (68) N-(2,6-diisopropylphenyl)-N'-(3-phenylpropyl)-N'-(4-phenylcyclohexyl)urea, melting at 196-197.5° C
- (69) N-(2,6-diisopropylphenyl)-N'-(2-phenylethyl)-N'-(4-phenylcyclohexyl)urea, melting at 157-159° C
- (70) cis-N-(2,6-diisopropylphenyl)-N'-benzyl-N'-(1,1'-bicyclohexyl-4-yl)urea, melting at 139° C
- (71) N-(2,4,6-trimethoxyphenyl)-N'-(4-phenylcyclohexyl)urea, melting at 166.5-168° C
- 50 (72) N-(2,6-diisopropylphenyl)-N'-cyclohexyl-N'-(4-phenylcyclohexyl)urea, melting at 58-62° C
- (73) cis-N-(2,6-diisopropylphenyl)-N'-(3-phenylpropyl)-N'-(4-phenylcyclohexyl)urea, melting at 147-149° C
- (74) trans-N-(2,6-diisopropylphenyl)-N'-(3-phenylpropyl)-N'-(4-phenylcyclohexyl)urea, melting at 195-196° C
- 55 (75) trans-N-(2,6-diisopropylphenyl)-N'-(1,1'-bicyclohexyl-4-yl)urea, melting at 283.5-284° C
- (76) trans-N-(2,6-diisopropylphenyl)-N'-benzyl-N'-(1,1'-bicyclohexyl-4-yl)urea, melting at 154-155° C
- (77) trans-N-(2,6-diisopropylphenyl)-N'-(1,3-benzodioxol-5-yl)methyl-N'-(1,1'-bicyclohexyl-4-yl)urea, melting at 147-148° C

- (78) trans-N-(2,6-diethylphenyl)-N'-cyclohexylmethyl-N'-(4-phenylcyclohexyl)urea, melting at 166-167° C
- (79) trans-N-(2,6-diisopropylphenyl)-N'-cyclohexylmethyl-N'-(4-phenylcyclohexyl)urea, melting at 150-152° C
- 5 (80) cis-N-(2,6-diisopropylphenyl)-N'-(3-chlorobenzyl)-N'-(1,1'-bicyclohexyl-4-yl)urea, melting at 174-176° C
- (81) cis-N-(2,6-diisopropylphenyl)-N'-(3-phenylpropyl)-N'-(1,1'-bicyclohexyl-4-yl)urea, melting at 148-149.5° C
- (82) trans-N-(2,6-diisopropylphenyl)-N'-(4-trifluoromethylbenzyl)-N'-(4-phenylcyclohexyl)urea 1/4hydrate, melting at 172-174° C
- 10 (83) N-(2,6-diisopropylphenyl)-N'-(2-morpholinoethyl)-N'-(4-phenylcyclohexyl)urea 1/4hydrate, melting at 174-176° C
- (84) N-(2,6-diisopropylphenyl)-N'-thenyl-N'-(4-phenylcyclohexyl)urea, melting at 150-152° C
- (85) cis-N-(2,6-diisopropylphenyl)-N'-(3,4,5-trimethoxybenzyl)-N'-(1,1'-bicyclohexyl-4-yl)urea, amorphous
- (86) cis-N-(2,6-diisopropylphenyl)-N'-(2-chloro-4-hydroxybenzyl)-N'-(1,1'-bicyclohexyl-4-yl)urea, melting at 168.5-170° C
- 15 (87) trans-N-(2,6-diisopropylphenyl)-N'-furfuryl-N'-(4-phenylcyclohexyl)urea, melting at 149-151° C
- (88) cis-N-(2,6-diisopropylphenyl)-N'-(4-chlorobenzyl)-N'-(1,1'-bicyclohexyl-4-yl)urea, melting at 160-161° C
- (89) cis-N-(2,6-diisopropylphenyl)-N'-(2-chlorobenzyl)-N'-(1,1'-bicyclohexyl-4-yl)urea, melting at 158-160° C
- 20 (90) N-(2,6-diethylphenyl)-N'-(2-methylbenzyl)-N'-(4-phenylcyclohexyl)urea, amorphous
- (91) N-(2,6-diethylphenyl)-N'-(2-chlorobenzyl)-N'-(4-phenylcyclohexyl)urea, melting at 150-151° C
- (92) trans-N-(2,6-diisopropylphenyl)-N'-(4-pyridylmethyl)-N'-(4-phenylcyclohexyl)urea 1/4hydrate, melting at 172-173° C
- 25 (93) trans-N-(2,6-diisopropylphenyl)-N'-(2-pyridylmethyl)-N'-(4-phenylcyclohexyl)urea hydrochloride 1/4hydrate, melting at 120-122° C
- (94) trans-N-(2,6-diisopropylphenyl)-N'-(2,3-difluorobenzyl)-N'-(4-phenylcyclohexyl)urea, melting at 168-170° C
- (95) trans-N-(2,6-diisopropylphenyl)-N'-(4-phenylbutyl)-N'-(4-phenylcyclohexyl)urea, melting at 151-152° C
- 30 (96) trans-N-(2,6-diisopropylphenyl)-N'-(2-ethoxybenzyl)-N'-(4-phenylcyclohexyl)urea, melting at 137-138° C
- (97) trans-N-(2,6-diisopropylphenyl)-N'-(2-bromobenzyl)-N'-(4-phenylcyclohexyl)urea, melting at 188-190° C
- 35 (98) trans-N-(2,6-diisopropylphenyl)-N'-(2-fluorobenzyl)-N'-(4-phenylcyclohexyl)urea, melting at 157-158° C
- (99) trans-N-(2,6-diisopropylphenyl)-N'-(2,3-dimethoxybenzyl)-N'-(4-phenylcyclohexyl)urea, melting at 160-162° C
- (100) N-(2,6-diisopropylphenyl)-N'-(2-methoxyethyl)-N'-(4-phenylcyclohexyl)urea, melting at 141-143° C
- 40 (101) trans-N-(2,6-diisopropylphenyl)-N'-(2,6-difluorobenzyl)-N'-(4-phenylcyclohexyl)urea, melting at 166-168° C
- (102) trans-N-(2,6-diisopropylphenyl)-N'-(2-hydroxybenzyl)-N'-(4-phenylcyclohexyl)urea, melting at 173-175° C
- (103) trans-N-(2,6-diisopropylphenyl)-N'-(2-(2-chlorophenyl)ethyl)-N'-(4-phenylcyclohexyl)urea, melting at 130-131° C
- 45 (104) trans-N-(2,6-diisopropylphenyl)-N'-(3-(2-chlorophenyl)propyl)-N'-(4-phenylcyclohexyl)urea, melting at 200-202° C
- (105) trans-N-(2,6-diisopropylphenyl)-N'-(2-methylthiobenzyl)-N'-(4-phenylcyclohexyl)urea
- (106) trans-N-(2,6-diisopropylphenyl)-N'-(2-benzoyloxybenzyl)-N'-(4-phenylcyclohexyl)urea, melting at 158.5-160° C
- 50 (107) trans-N-(2,6-diisopropylphenyl)-N'-(2-(2-methylphenyl)ethyl)-N'-(4-phenylcyclohexyl)urea, melting at 122-123° C
- (108) trans-N-(2,6-diisopropylphenyl)-N'-(2-benzylthiobenzyl)-N'-(4-phenylcyclohexyl)urea
- (109) trans-N-(2,4,6-tri-tert-butylphenyl)-N'-benzyl-N'-(4-phenylcyclohexyl)urea
- 55 (110) trans-N-(2,6-diisopropylphenyl)-N'-(3-(2-methylphenyl)propyl)-N'-(4-phenylcyclohexyl)urea
- (111) trans-N-(2,6-diisopropylphenyl)-N'-(4-(2-chlorophenyl)butyl)-N'-(4-phenylcyclohexyl)urea
- (112) trans-N-(2,6-diethylphenyl)-N'-(2-(2-methylphenyl)ethyl)-N'-(4-phenylcyclohexyl)urea
- (113) trans-N-(2,6-diisopropylphenyl)-N'-(4-hydroxy-3,5-di-tert-butylbenzyl)-N'-(4-phenylcyclohexyl)urea

- (114) trans-N-(2,6-diisopropylphenyl)-N'-(2-(2-chlorophenoxy)ethyl)-N'-(4-phenylcyclohexyl)urea
 (115) trans-N-(2,6-dimethylphenyl)-N'-benzyl-N'-(4-phenylcyclohexyl)urea, melting at 119-121 ° C
 (116) trans-N-(2,6-diisopropylphenyl)-N'-(4-(2-methylphenyl)butyl)-N'-(4-phenylcyclohexyl)urea
 (117) trans-N-(2,6-diethylphenyl)-N'-(2-(2-chlorophenyl)ethyl)-N'-(4-phenylcyclohexyl)urea
 5 (118) trans-N-(2,4,6-trimethoxyphenyl)-N'-(2-(2-methylphenyl)ethyl)-N'-(4-phenylcyclohexyl)urea
 (119) trans-N-(2,6-diisopropylphenyl)-N'-(2-ethylbenzyl)-N'-(4-phenylcyclohexyl)urea
 (120) cis-N-(2,6-diisopropylphenyl)-N'-(2-methoxybenzyl)-N'-(4-phenylcyclohexyl)urea, melting at 145-146.5 ° C

10 Example of Pharmaceutical Composition

The tablets containing the compound of formula (I) can be prepared by the following formulation.

15	Compound (I)	100 mg
	Lactose	76 mg
	Corn starch	10 mg
	Carboxymethylcellulose calcium	5 mg
	Methylcellulose	3 mg
20	Magnesium stearate	2 mg
	Polyvinyl pyrrolidone	4 mg
	Total	200 mg

Compound (I) is crushed with an atomizer to make fine powder having an average particle size below
 25 10 μ . The fine powder of Compound (I), lactose, corn starch, carboxymethylcellulose calcium and methylcellulose are mixed well in a kneader and then kneaded with a binder prepared by polyvinyl pyrrolidone. The wet mass is passed through a 200 mesh sieve to give granules and then dried in an oven at 50 ° C. The dry granules containing 3-4% of water content is forced through a 24 mesh sieve. Magnesium stearate is mixed with and compressed into tablets by using a rotatory tableting machine with a flat punch of 8 mm diameter.

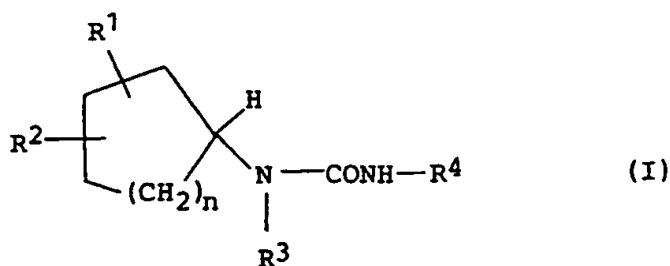
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Claims

1. A cycloalkylurea compound of the formula (I):

35

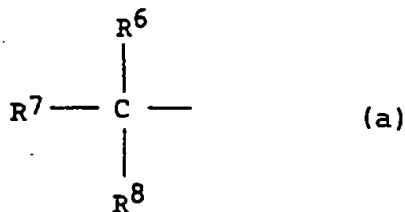
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wherein R¹ is a group of the formula (a):

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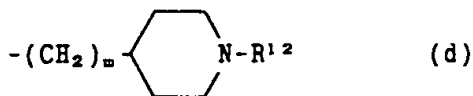
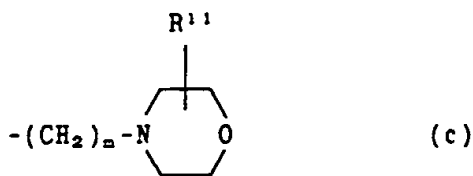
(wherein R⁶, R⁷ and R⁸ are the same or different and each is hydrogen or alkyl having 1 to 8 carbon atoms, with the proviso that compounds wherein two or three of R⁶, R⁷ and R⁸ are hydrogens are

excluded), haloalkyl, cycloalkyl, alkoxy having 1 to 5 carbon atoms, phenyl, aralkyl or heteroaryl, or phenyl, aralkyl or heteroaryl each substituted on the aromatic ring by 1 to 3 substituents selected from alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 8 carbon atoms, cycloalkyl, cycloalkyloxy, haloalkyl, haloalkoxy, halogen, nitro, amino and substituted amino;

R^2 is hydrogen, phenyl or phenyl substituted by 1 to 3 substituents selected from alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, haloalkyl and halogen, whererin R^1 and R^2 may be substituted on the same carbon atom of cycloalkyl ring;

R^1 and R^2 together may form a substituted or unsubstituted cyclic hydrocarbon having 3 to 7 carbon atoms or a substituted or unsubstituted spiran;

R^3 is hydrogen, alkyl having 1 to 8 carbon atoms, cycloalkyl, cycloalkylalkyl, alkoxyalkyl, alkylthioalkyl, aralkyl substituted by alkylendioxy, aralkyl, aralkyloxyalkyl, aralkyl or aralkyloxyalkyl each substituted on the aromatic ring by 1 to 3 substituents selected from alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 5 carbon atoms, halogen, nitro, hydroxy, amino, substituted amino, haloalkyl, alkylthio, benzyloxy and benzylthio, heteroarylalkyl, phenoxyalkyl, or heteroarylalkyl or phenoxyalkyl each substituted on the aromatic ring by 1 to 3 substituents selected from halogen, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, nitro, amino and haloalkyl, or a group of the formula (c) or (d):



wherein m is 1 or 2, R^{11} is hydrogen or alkyl having 1 to 4 carbon atoms, R^{12} is hydrogen, alkyl having 1 to 4 carbon atoms or aralkyl;

R^4 is phenyl substituted by 1 to 3 substituents selected from alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, haloalkyl, halogen, amino and substituted amino;

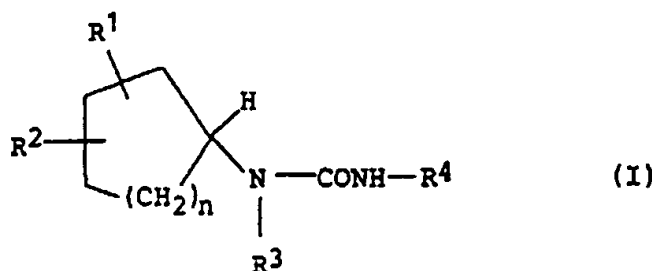
n is 1, 2 or 3;

or a pharmaceutically acceptable salt thereof.

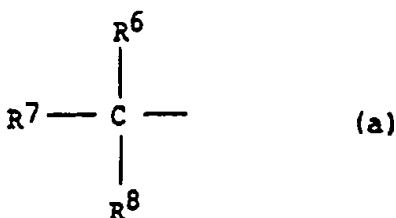
- 40 2. A compound as claimed in Claim 1 wherein R^1 is phenyl or phenyl substituted by 1 to 3 substituents selected from alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 8 carbon atoms, haloalkyl, halogen, nitro, amino and substituted amino; R^2 is phenyl or phenyl substituted by 1 to 3 substituents selected from alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, trifluoromethyl and halogen, whererin R^1 and R^2 may be substituted on the same carbon atom of cycloalkyl ring; or R^1 and R^2
- 45 together may form a substituted or unsubstituted cyclic hydrocarbon having 3 to 7 carbon atoms or a substituted or unsubstituted spiran; R^3 is hydrogen, alkyl having 1 to 8 carbon atoms, cycloalkyl, cycloalkylalkyl, aralkyl substituted by alkylendioxy, aralkyl, aralkyl substituted on the aromatic ring by 1 to 3 substituents selected from alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 5 carbon atoms, halogen, nitro, hydroxy, haloalkyl, alkylthio, benzyloxy and benzylthio, or heteroarylalkyl; R^4 is phenyl substituted by 1 to 3 substituents selected from alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, trifluoromethyl, halogen and substituted amino; n is 2;
- 50 or a pharmaceutically acceptable salt thereof.
3. A compound as claimed in Claim 1 wherein R^1 is secondary or tertiary alkyl having 3 to 5 carbon atoms, cycloalkyl or phenyl; R^2 is hydrogen; R^3 is hydrogen, alkyl having 1 to 8 carbon atoms, cycloalkyl, cycloalkylalkyl, aralkyl substituted by alkylendioxy, aralkyl, aralkyl substituted on the aromatic ring by 1 to 3 substituents selected from alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 5 carbon atoms, halogen, nitro, hydroxy, haloalkyl, alkylthio, benzyloxy and benzylthio, or heteroarylal-
- 55

kyl; R⁴ is phenyl substituted by 1 to 3 substituents selected from alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms and halogen; n is 2; or a pharmaceutically acceptable salt thereof.

4. A compound as claimed in Claim 1 selected from the group consisting of
- 5 N-(2,6-diisopropylphenyl)-N'-(2-phenylcyclohexyl)urea,
 N-(2,6-diisopropylphenyl)-N'-heptyl-N'-(4-phenylcyclohexyl)urea,
 trans-N-(2,6-diisopropylphenyl)-N'-benzyl-N'-(4-phenylcyclohexyl)urea,
 N-(2,6-diethylphenyl)-N'-benzyl-N'-(4-phenylcyclohexyl)urea,
 N-(2,4-difluorophenyl)-N'-benzyl-N'-(4-phenylcyclohexyl)urea,
 10 N-(2,6-diisopropylphenyl)-N'-(3,4-dimethoxybenzyl)-N'-(4-phenylcyclohexyl)urea,
 N-(2,4,6-trimethoxyphenyl)-N'-benzyl-N'-(4-phenylcyclohexyl)urea,
 N-(2,6-diisopropylphenyl)-N'-benzyl-N'-(4-tert-butylcyclohexyl)urea,
 trans-N-(2,6-diisopropylphenyl)-N'-(2-methylbenzyl)-N'-(4-phenylcyclohexyl)urea,
 trans-N-(2,6-diisopropylphenyl)-N'-(2-chlorobenzyl)-N'-(4-phenylcyclohexyl)urea,
 15 N-(2,6-diisopropylphenyl)-N'-(2-phenylethyl)-N'-(4-phenylcyclohexyl)urea,
 trans-N-(2,6-diisopropylphenyl)-N'-(3-phenylpropyl)-N'-(4-phenylcyclohexyl)urea,
 trans-N-(2,6-diisopropylphenyl)-N'-(1,1'-bicyclohexyl-4-yl)urea,
 trans-N-(2,6-diethylphenyl)-N'-cyclohexylmethyl-N'-(4-phenylcyclohexyl)urea,
 trans-N-(2,6-diisopropylphenyl)-N'-cyclohexylmethyl-N'-(4-phenylcyclohexyl)urea,
 20 N-(2,6-diisopropylphenyl)-N'-thenyl-N'-(4-phenylcyclohexyl)urea,
 trans-N-(2,6-diisopropylphenyl)-N'-furfuryl-N'-(4-phenylcyclohexyl)urea,
 N-(2,6-diethylphenyl)-N'-(2-methylbenzyl)-N'-(4-phenylcyclohexyl)urea,
 N-(2,6-diethylphenyl)-N'-(2-chlorobenzyl)-N'-(4-phenylcyclohexyl)urea and
 trans-N-(2,6-diisopropylphenyl)-N'-(2-pyridylmethyl)-N'-(4-phenylcyclohexyl)urea
 25 or a pharmaceutically acceptable salt thereof.
5. A compound as claimed in Claim 1 selected from the group consisting of
- trans-N-(2,6-diisopropylphenyl)-N'-benzyl-N'-(4-phenylcyclohexyl)urea,
 trans-N-(2,6-diisopropylphenyl)-N'-(2-methylbenzyl)-N'-(4-phenylcyclohexyl)urea,
 30 trans-N-(2,6-diisopropylphenyl)-N'-(2-chlorobenzyl)-N'-(4-phenylcyclohexyl)urea and
 trans-N-(2,6-diethylphenyl)-N'-cyclohexylmethyl-N'-(4-phenylcyclohexyl)urea,
 or a pharmaceutically acceptable salt thereof.
6. A compound as claimed in Claim 1 which is N-(2,6-diisopropylphenyl)-N'-(4,4-diphenylcyclohexyl)urea,
 35 or a pharmaceutically acceptable salt thereof.
7. A pharmaceutical composition consisting of an effective amount of compound as claimed in Claim 1 or
 a pharmaceutically acceptable salt thereof and pharmaceutical carriers.
- 40 8. Process for preparing a cycloalkylurea compound of the formula (I):



55 wherein R¹ is a group of the formula (a):

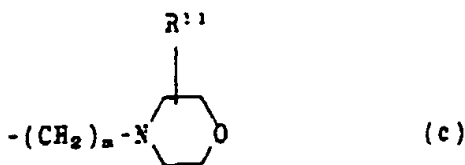


(wherein R^6 , R^7 and R^8 are the same or different and each is hydrogen or alkyl having 1 to 8 carbon atoms, with the proviso that compounds wherein two or three of R^6 , R^7 and R^8 are hydrogens are excluded), haloalkyl, cycloalkyl, alkoxy having 1 to 5 carbon atoms, phenyl, aralkyl or heteroaryl, or phenyl, aralkyl or heteroaryl each substituted on the aromatic ring by 1 to 3 substituents selected from alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 8 carbon atoms, cycloalkyl, cycloalkoxy, haloalkyl, haloalkoxy, halogen, nitro, amino and substituted amino;

R^2 is hydrogen, phenyl or phenyl substituted by 1 to 3 substituents selected from alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, haloalkyl and halogen, wherein R^1 and R^2 may be substituted on the same carbon atom of cycloalkyl ring;

R^1 and R^2 together may form a substituted or unsubstituted cyclic hydrocarbon having 3 to 7 carbon atoms or a substituted or unsubstituted spiran;

R^3 is hydrogen, alkyl having 1 to 8 carbon atoms, cycloalkyl, cycloalkylalkyl, alkoxyalkyl, alkylthioalkyl, aralkyl substituted by alkylenedioxy, aralkyl, aralkyloxyalkyl, aralkyl or aralkyloxyalkyl each substituted on the aromatic ring by 1 to 3 substituents selected from alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 5 carbon atoms, halogen, nitro, hydroxy, amino, substituted amino, haloalkyl, alkylthio, benzyloxy and benzylthio, heteroarylalkyl, phenoxyalkyl, or heteroarylalkyl or phenoxyalkyl each substituted on the aromatic ring by 1 to 3 substituents selected from halogen, alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, nitro, amino and haloalkyl, or a group of the formula (c) or (d):



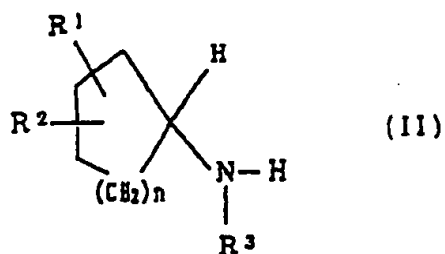
wherein m is 1 or 2, R^{11} is hydrogen or alkyl having 1 to 4 carbon atoms, R^{12} is hydrogen, alkyl having 1 to 4 carbon atoms or aralkyl;

R^4 is phenyl substituted by 1 to 3 substituents selected from alkyl having 1 to 4 carbon atoms, alkoxy having 1 to 4 carbon atoms, haloalkyl, halogen, amino and substituted amino;

n is 1, 2 or 3;

or a pharmaceutically acceptable salt thereof, which comprises:

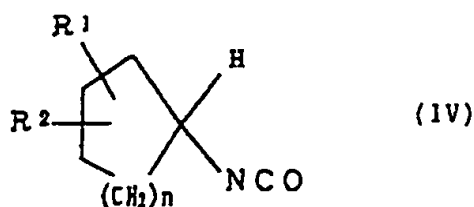
a) reacting a cycloalkylamine of the formula (II):



wherein each symbol is as defined above,
or a salt thereof with an isocyanic ester of the formula (III):



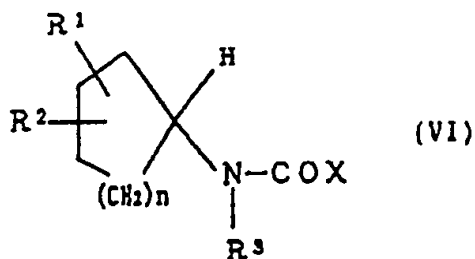
wherein R⁴ is as defined above,
in an inert solvent at temperatures between -20° C and 150° C, or
b) for preparing a compound of the formula (I) wherein R³ is hydrogen, reacting an isocyanic ester of
20 the formula (IV):



wherein each symbol is as defined above, with an amine of the formula (V):



wherein R⁴ is as defined above,
in an inert solvent at temperatures between -20° C and 150° C, or
c) for preparing a compound of the formula (I) wherein R³ is the substituent other than hydrogen,
40 reacting a carbonyl halide of the formula (VI):



wherein X is halogen and other symbols are as defined above, with an amine of the formula (V):



wherein R⁴ is as defined above,
in an inert solvent at temperatures between -20° C and 150° C.



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PARTIAL EUROPEAN SEARCH REPORT

Application Number

which under Rule 45 of the European Patent Convention shall be considered, for the purposes of subsequent proceedings, as the European search report

EP 92 10 5655

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl.5)
D,Y	US-A-4 387 105 (V.G. DeVRIES et al.) * Whole document, especially column 6, table I, line 45 *	1,3,4,7	C 07 C 275/28 A 61 K 31/17 C 07 C 275/30
Y	EP-A-0 344 425 (WARNER-LAMBERT CO.) * Pages 9-10, claims 1-5; pages 7-8, examples 6,8 *	1,3,4,7	C 07 C 275/40 C 07 D 213/40 C 07 D 295/13 C 07 D 307/52
A	EP-A-0 335 375 (WARNER-LAMBERT CO.) * Pages 16-18, claims 1-5; page 14, example 9 *	1,3,4,7	C 07 D 317/58 C 07 D 333/20
D,A	EP-A-0 384 320 (WARNER-LAMBERT CO.) * Pages 14-16, claims 1-5,7-9,11-12,15; page 10, example 2; pages 12-13, example 9 *	1-7	
			TECHNICAL FIELDS SEARCHED (Int. Cl.5)
			C 07 C
INCOMPLETE SEARCH			
<p>The Search Division considers that the present European patent application does not comply with the provisions of the European Patent Convention to such an extent that it is not possible to carry out a meaningful search into the state of the art on the basis of some of the claims</p> <p>Claims searched completely : 2-8 Claims searched incompletely : 1 Claims not searched :</p> <p>Reason for the limitation of the search: Lack of conciseness (Art.84 EPC)</p> <p>Claim 1 is too broad with respect to the definition of the substituents R¹,R² (their flexible attachment points), R³ and the ringsize (CH₂)_n. Hence, the full range of claim 1 cannot be searched comprehensively, not even by means of on-line methods.</p>			
Place of search	Date of completion of the search	Examiner	
THE HAGUE	10-07-1992	FINK D.	
CATEGORY OF CITED DOCUMENTS		<p>T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons</p> <p>Δ : member of the same patent family, corresponding document</p>	
<p>X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document</p>			

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